California Medical Device Recall Information

Recall Name
Hospira Recalls Plum A+ and Plum A+3 Infusion Systems Due to Potential Alarm Volume Failure

Recall Date	Product Description	Recalling Firm	Recall Reason
5/28/14	Infusion Pumps: **Plum A+** List Numbers: • 11005 (Plum Hyperbaric) • 11971 • 12391 • 20679 • 20792 **Plum A+3** List Numbers • 12618 • 20678	**Hospira, Inc.** Lake Forest, IL	*Some of the alarms may fail to sound in situations that should trigger a sound.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	Affected Lot Numbers Affected Lot Numbers for List Number 11005	**CA**, nationwide	Distributed: July 2012 to May 2014

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm436770.htm